

Lower Ferry Road, Ewing, New Jersey 08618–1414 applied to be registered as a bulk manufacturer of the following basic classes controlled substances:

Controlled substance	Drug code	Schedule
Pentobarbital	2270	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Remifentanyl	9739	II
Fentanyl	9801	II

The company plans to initially to manufacture API quantities of the listed controlled substances for validation purposes and FDA approval, then eventually up FDA approval to produce commercial size batches for distribution to dosage form manufacturers.

Dated: December 19, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31282 Filed 12–27–16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement

Administration (DEA) as bulk manufacturers of various classes of controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as manufacturers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted for these notices.

Company	FR docket	Published
National Center for Natural Products Research (NIDA MPROJECT)	80 FR 78766	December 17, 2015.
Cambrex Charles City	81 FR 46956	July 19, 2016.
Chemtos, LLC	81 FR 54604	August 16, 2016.
Stepan Company	81 FR 54849	August 17, 2016.
Chattem Chemicals	81 FR 57932	August 24, 2016.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company’s maintenance of effective controls against diversion by inspecting and testing each company’s physical security systems, verifying each company’s compliance with state and local laws, and reviewing each company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed persons.

Dated: December 19, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31280 Filed 12–27–16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Synthcon, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before February 27, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated her authority under the Controlled

Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on December 16, 2015, Synthcon, LLC, 770 Wooten Road, Unit 101, Colorado Springs, Colorado 80915 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
3-Fluoro-N-methylcathinone (3-FMC)	1233	I
Cathinone	1235	I
Methcathinone	1237	I
4-Fluoro-N-methylcathinone (4-FMC)	1238	I
Pentedrone (α-methylaminovalerophenone)	1246	I